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| APPLICATION NO. | FILING DATE | FIRST NAMED INVENTOR | ATTORNEY DOCKET NO. | CONFIRMATION NO. |
|-----------------|-------------|----------------------|---------------------|------------------|
| 10/753,289 | 01/05/2004 | Steven M. Watkins | 475512000100 | 1703 |

25226 7590 06/19/2007
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| EXAMINER |
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NEGIN, RUSSELL SCOTT

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| ART UNIT | PAPER NUMBER |
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1631

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| MAIL DATE | DELIVERY MODE |
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06/19/2007

PAPER

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

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|------------------------------|------------------------|--|---------------------|--|
| Office Action Summary | Application No. | | Applicant(s) | |
| | 10/753,289 | | WATKINS, STEVEN M. | |
| | Examiner | | Art Unit | |
| | Russell S. Negin | | 1631 | |

-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --

Period for Reply

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) OR THIRTY (30) DAYS, WHICHEVER IS LONGER, FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

Status

- 1) ☐ Responsive to communication(s) filed on 19 March 2007.
- 2a) ☐ This action is **FINAL**. 2b) ☒ This action is non-final.
- 3) ☐ Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

Disposition of Claims

- 4) ☒ Claim(s) 2-17 and 56-58 is/are pending in the application.
- 4a) Of the above claim(s) _____ is/are withdrawn from consideration.
- 5) ☐ Claim(s) _____ is/are allowed.
- 6) ☒ Claim(s) 2-17 and 56-58 is/are rejected.
- 7) ☐ Claim(s) _____ is/are objected to.
- 8) ☐ Claim(s) _____ are subject to restriction and/or election requirement.

Application Papers

- 9) ☐ The specification is objected to by the Examiner.
- 10) ☐ The drawing(s) filed on _____ is/are: a) ☐ accepted or b) ☐ objected to by the Examiner.
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).
Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).
- 11) ☐ The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.

Priority under 35 U.S.C. § 119

- 12) ☐ Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).
- a) ☐ All b) ☐ Some * c) ☐ None of:
1. ☐ Certified copies of the priority documents have been received.
 2. ☐ Certified copies of the priority documents have been received in Application No. _____.
 3. ☐ Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).

* See the attached detailed Office action for a list of the certified copies not received.

Attachment(s)

- | | |
|--|---|
| 1) <input type="checkbox"/> Notice of References Cited (PTO-892) | 4) <input type="checkbox"/> Interview Summary (PTO-413) Paper No(s)/Mail Date. _____ |
| 2) <input type="checkbox"/> Notice of Draftsperson's Patent Drawing Review (PTO-948) | 5) <input type="checkbox"/> Notice of Informal Patent Application |
| 3) <input checked="" type="checkbox"/> Information Disclosure Statement(s) (PTO/SB/08) Paper No(s)/Mail Date <u>3/29/06</u> | 6) <input type="checkbox"/> Other: _____ |

DETAILED ACTION

Comments

Applicants' amendments and request for reconsideration in the communication filed on 19 March 2007 are acknowledged and the amendments are entered.

Claims 2-17 and 56-58 are pending and examined in the instant Office action.

Information Disclosure Statement

The documents missing from the information Disclosure Statement filed 29 March 2006 have been provided. A corrected copy of the signed Information disclosure statement is provided with this Office action.

Claim Rejections - 35 USC § 102

The following is a quotation of the appropriate paragraphs of 35 U.S.C. 102 that form the basis for the rejections under this section made in this Office action:

A person shall be entitled to a patent unless –

(b) the invention was patented or described in a printed publication in this or a foreign country or in public use or on sale in this country, more than one year prior to the date of application for patent in the United States.

The rejections of claims 2, 3, 5, 6, 12, and 54 under 35 U.S.C. 102(b) as being anticipated by Wong et al. [Magnetic Resonance in Medicine, 1994, volume 32, page 440-446] are withdrawn in view of amendments made by applicant to the set of claims filed on 19 March 2007.

The rejection of claims 2, 3, 12, 55, and 56 under 35 U.S.C. 102(b) as being anticipated by Ruan et al. [Journal of Dairy Science, volume 81, 1998, pages 9-15] in light of the definition of "Heat map" in Wikipedia [accessed at http://en.wikipedia.org/wiki/Heat_map on 6 December 2006] are withdrawn in view of amendments made by applicant to the set of claims filed on 19 March 2007.

The following rejection is reiterated from the Office action of 18 December 2006 and is newly applied to claims 5, 6, and 57-58:

Claims 2, 3, 5, 6, 12, and 56-58 are rejected under 35 U.S.C. 102(b) as being anticipated by Ruan et al. [Journal of Dairy Science, volume 81, 1998, pages 9-15].

Claim 2 is drawn to a method for presenting analysis of a plurality of quantitative lipid profiles comprising designating the plurality of quantitative lipid profiles, identifying differences or similarities in the plurality of lipid metabolites between the quantitative lipid profiles, and displaying the identified differences or similarities on a heat map.

The study of Ruan et al., entitled, "A magnetic resonance imaging technique for quantitative mapping of moisture and fat in a cheese block," is drawn to analysis of water and oil suspensions, as stated in the abstract:

Separate magnetic resonance images of water fat of oil-in-water emulsions and cheese blocks were obtained using the chemical shift selective suppression technique. With this technique, the proton signals emitted from water can be readily separated from those emitted from fat in the same sample through a single experiment using magnetic resonance imaging. Relaxation compensation was made to improve the quality of suppression. The experiment using oil-in-water emulsions demonstrated an excellent linear relationship between the intensity of the signal and the concentration of water or fat.

The Materials and Methods section on the second column of page 10 of Ruan et al. elaborates on the procedures of the method described in the abstract:

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Oil-in water-emulsions. Oil in water emulsions were prepared to serve as homogeneous samples for the purpose of development and testing of the MRI techniques. The emulsions were freshly made before the MRI experiments by taking known amounts of vegetable oil (Crisco oil; Proctor & Gamble, Cincinnati, OH) with 3 mM CuSO₄ in 20 mm diameter glass tubes... Percentages of oil by volume were 0, 20, 30, 40, 50, 60, 70, and 100%. Three percent of Tween 40 (polyoxyethylene sorbitan monooleate) by volume of the oil was added to improve the stability of the mixture.

Once these oil and water emulsions are made, they are used to generate multiple "lipid metabolite profiles" as explained below.

To clarify the definition of "lipid metabolite profiles," metabolites are defined on page 8 of the instant specification as:

A biomolecule that has a functional and/or compositional role (such as a component of a membrane) in a biological system, and which is not a molecule of DNA, RNA, or protein. Examples of metabolites include lipids, carbohydrates, vitamins, co-factors, pigments, and so forth. Metabolites can be obtained through the diet (consumed from the environment) or synthesized within an organism... By profiling the metabolite composition of a biological sample, for instance using the methods described herein, data on genotype, metabolism, and diet can be obtained in great detail.

Consequently, the language of this definition does not eliminate the possibility that any molecule in a biological system that has a functional or compositional role (that is not DNA, RNA, or protein) is a metabolite. For example, both water and oils fit this definition in the instant specification.

Metabolite profiles are defined in the instant specification on page 9, lines 3-5, as:

The set of data produced from analysis of an individual sample is referred to herein as a individual [sic] lipid profile/metabolic profile ("lipomic profile") of that sample. Certain examples of lipid metabolite profiles include a highly comprehensive set of metabolic measurements (a profile) by multi-parallel analyses.

The comparison of two metabolite profiles of similar scope (i.e. containing information about the same or a similar or overlapping set or subset of metabolites) from cells/tissues/subjects that have been differently treated, or that are genetically different based on disease state or condition, provides information on the metabolic effects of the difference.

As stated in the claims cited above, the language of the instant claims uses “lipid metabolic profiles,” which narrows the scope of the invention to metabolic profiles of lipids.

In the section “Imaging and quantification of water and fat in homogenous fat and water phantoms” on page 12 of Ruan et al., Ruan et al. finds two linear equations corresponding to Figures 5A and 5B, respectively, each of which is interpreted as a “lipid metabolic profile.”

Figure 5A is the lipid metabolite profile of the oil-in-water emulsions in a water suppressed MRI image while Figure 5B is the lipid metabolite profile of the same set of oil in water emulsions in a fat suppressed image. Equations 1 and 2 quantify the intensities of the lipid metabolite profiles in Figures 5A and 5B, respectively, to reveal the similarity that the two correlations between the concentration of oil (and water) in the binary mixture and image intensity are linear. The cited definitions and claims do not require each “lipid metabolite profile” to be taken from different sets of samples.

In Figure 5B, the concentration of water is measured because the lipids are suppressed in the image. Water is not a lipid, but in a binary mixture of water and oil, the concentration of water is directly and uniquely dependent on the amount of oil added (the volume of the mixture not occupied by water can only be occupied by oil).

Ruan et al. elaborates on this generation of a plurality metabolite profiles generated from the series of oil in water emulsions described above on page 12, column 2:

Figure 5 shows a series of water suppressed (A) and fat suppressed (B) magnetic resonance images of the oil-in-water emulsion phantoms. Greater brightness indicates a stronger signal. The signal intensities vary as the water and fat contents vary. When signals that were emitted

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from water were suppressed (Figure 5A), the signal intensity increased as water content increased. When fat or water was absent from the mixture, little signal can be seen in the water-suppressed or fat-suppressed images, indicating that reliable selective signal suppression was achieved... An excellent linear correlation between the MRI signal intensity and water and fat contents was found.

Figure 5 on page 13 of Ruan et al. illustrates a "heat map" of the fat contents of a variety of different lipid composition profiles. The caption to Figure 5 states, "Magnetic resonance images of oil-in-water phantoms: A. water-suppressed images (the percentages indicate the oil contents of the mixtures); B. oil-suppressed images (the percentages indicate the water contents of the mixtures)."

Consequently, Ruan et al. shows analysis of multiple lipid mixtures using magnetic resonance imaging, which are designated and mapped in Figure 5 of Ruan et al.

The term "heat map" is not explicitly recited in Ruan et al.

On page 37, lines 17-19, applicant defines heat map as:

In a heat map display, quantitative metabolite data from a test sample is compared to quantitative metabolite data from a base line or standard sample (a control) and the increase or decrease in each metabolite is indicated on the display, usually in a readily recognizable fashion.

It is inherent that the illustration in Figure 5 of Ruan et al. is a heat map because it is a two dimensional map of multiple lipid profiles marked by shades of colors.

Claim 3 is dependent from claim 2 with the additional limitation of the quantitative lipid metabolite profiles comprising quantitative measures of at least two lipids wherein the quantified measurements are obtained using an internal standard for at least one of the lipids.

Figure 5 on page 13 of Ruan et al. illustrates a “heat map” of the fat contents of a variety of different lipid composition profiles. The caption to Figure 5 states, “Magnetic resonance images of oil-in-water phantoms: A. water-suppressed images (the percentages indicate the oil contents of the mixtures); B. oil-suppressed images (the percentages indicate the water contents of the mixtures).”

Consequently, Ruan et al. shows analysis of multiple lipid mixtures using magnetic resonance imaging, which are designated and mapped in Figure 5 of Ruan et al.

It is inherent that a sample of Crisco vegetable oil comprises a mixture of multiple lipids that is compared to the standards of 0% lipid content and 100% lipid content in Figure 5 of Ruan et al.

Claim 5 is dependent from claim 2 with the additional limitation that the quantitative lipid metabolite profiles each comprise a quantified measurement of a lipid in a lipid class.

Claim 6 is dependent from claim 5, wherein the quantified measurement of the lipid in the lipid class is obtained using an internal standard for the lipid class.

Figure 5 of Ruan et al. illustrates the standards at the 0 percent and 100 percent endpoints of Figure in which the mixture is either entirely oil or water to which the linear correlations in the study are applied.

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Claim 12 is dependent from claim 2 wherein at least one of the quantitative lipid metabolite profiles is generated comprising separating a biological sample into fractions based on a plurality of lipid classes, wherein at least one quantitative internal standard is included for each lipid class; and measuring the quantity of a plurality of lipid metabolites in the fractions.

Again, Figure 5 of Ruan et al. illustrates the biological samples separated into fractions where water is the standard against which the lipid classes are measured.

Claim 56 is dependent from claim 2 wherein an increase or decrease in the lipid metabolite is indicated on the heat map by a color and the relevant amount of the increase or decrease is indicated by the intensity of the color.

The "heat maps" in Figure 5 of Ruan et al. illustrate such a trend in colors with respect to lipid content.

Claim 57 is dependent from claim 2, further comprising generating a written report.

The figures, tables and equations of Ruan et al. serve as written reports.

Claim 58 is dependent from claim 2 wherein one of the quantitative lipid metabolite profiles is a control lipomic profile.

In this instance, Figure 5 of Ruan et al. shows the control lipomic profiles at the 100% and 0% levels of each lipid/water.

Applicant's arguments filed 19 March 2007 have been fully considered but they are not persuasive. New grounds for rejection are applied necessitated by the amendment of the applicants.

Applicant first argues on page 9 of the Remarks of 19 March 2007 that the definition of "heat map" in the previous Office action is inconsistent with the definition employed in the specification. On page 37, lines 17-19, applicant defines heat map as:

In a heat map display, quantitative metabolite data from a test sample is compared to quantitative metabolite data from a base line or standard sample (a control) and the increase or decrease in each metabolite is indicated on the display, usually in a readily recognizable fashion.

This revised definition is considered in the instant Office action, but does not eliminate the existence of prior art rejections.

Applicant next argues on page 10 of the Remarks:

...the Ruan et al. reference does not compare the data in Figure 5A to the data in Figure 5B to identify and display the differences or similarities between the two sets of quantitative data in the form of a heat map.

This argument is not found to be persuasive because the lipid profiles have their intensities compared by correlations (i.e. "r" values) in equation 1 on page 12 of Ruan et al. for Figure 5A and equation 2 on page 12 of Ruan et al. for Figure 5B. According to the results of the equations, Figure 5B shows a slightly more linear correlation ($r^2 = 0.97961$) than Figure 5A ($r^2 = 0.9764$).

Applicant next argues on page 10 of the Remarks that the analysis in Figure 5 does not present a plurality of lipid metabolites. This argument is not found to be

persuasive because Crisco is inherently a mixture of many lipids found in vegetables (i.e. vegetable oil is inherently not composed of a single oil or lipid).

Since the claims are amended, the 35 U.S.C. 102 rejection involving Wong et al. is no longer applicable.

Claim Rejections - 35 USC § 103

The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negated by the manner in which the invention was made.

The rejections of claims 2 and 57 under 35 U.S.C. 103(a) as being unpatentable over Wong et al. in view of Moser et al. [Neurochemical Research, volume 24, 1999, pages 187-197] are withdrawn in view of amendments made by applicant to the set of claims filed on 19 March 2007.

The rejections of claims 2, 12, 15, and 16 under 35 U.S.C. 103(a) as being unpatentable over Wong et al. in view of Moser et al. [Moser et al., Annals of Neurology, volume 45, 1999, pages 100-110] are withdrawn in view of amendments made by applicant to the set of claims filed on 19 March 2007.

The rejections of claims 2, 3, 6, 7, 9, 12 and 13 under 35 U.S.C. 103(a) as being unpatentable over Wong et al. in view of Watkins et al. [Journal of Lipid Research, volume 39, 1998, pages 1583-1588] are withdrawn in view of amendments made by applicant to the set of claims filed on 19 March 2007.

The rejections of claims 2-4, 12, and 14 under 35 U.S.C. 103(a) as being unpatentable over Wong et al. in view of Watkins et al. and in further view of Siguel [US Patent 5,075,101; IDS of 1/5/2004] are withdrawn in view of amendments made by applicant to the set of claims filed on 19 March 2007.

The rejections of claims 2, 17, and 19 under 35 U.S.C. 103(a) as being unpatentable over Wong et al. in view of "The World of Membrane Lipids," [www.biochem.Missouri.edu/~lesa/LIPIDS/membrane_lipid.html; accessed on 6 December 2006, page made on 2 February 1999] are withdrawn in view of amendments made by applicant to the set of claims filed on 19 March 2007.

The rejections of claims 2, 17, and 18 under 35 U.S.C. 103(a) as being unpatentable over Ruan et al. in view of "The World of Membrane Lipids," [www.biochem.Missouri.edu/~lesa/LIPIDS/membrane_lipid.html; accessed on 6 December 2006, page made on 2 February 1999] are withdrawn in view of amendments made by applicant to the set of claims filed on 19 March 2007.

The rejections of claims 2 and 10 under 35 U.S.C. 103(a) as being unpatentable over Wong et al. in view of Grav et al. [Journal of Chromatography B, volume 658, 1994, pages 1-10] are withdrawn in view of amendments made by applicant to the set of claims filed on 19 March 2007.

The rejections of claims 2, 5, and 8 under 35 U.S.C. 103(a) as being unpatentable over Wong et al. in view of Dutta et al. [JAOCS, volume 74, no. 6, 1997, pages 647-657] are withdrawn in view of amendments made by applicant to the set of claims filed on 19 March 2007.

The following rejections are newly applied:

Rejection #1 under 35 U.S.C. 103(a):

Claims 15 and 16 are rejected under 35 U.S.C. 103(a) as being unpatentable over Ruan et al. as applied to claims 2, 3, 5, 6, 12, and 56-58 above in further view of Moser et al. [Moser et al., *Annals of Neurology*, volume 45, 1999, pages 100-110].

Claims 15 and 16 depend from claim 2 wherein the separating and measuring methods comprise chromatography.

Ruan et al. as applied to claims 2, 3, 5, 6, 12, and 56-58 above does not teach chromatography.

The study of Moser et al., entitled, "Plasma very long chain fatty acids in 3,000 peroxisome disease patients and 29,000 controls," states in the first sentence of the abstract, "The assay of plasma very long chain fatty acids (VLCFAs), developed in our laboratory on 1981, has become the most widely used procedure for the diagnosis of X-linked adrenoleukodystrophy (X-ALD) and other peroxisomal disorders."

The second column of page 101 of Moser et al. (*Neurology*) states:

Capillary Gas Liquid Chromatographic Analysis of Very Long Chain Fatty Acids
The VLCFA assay procedure was described in 1981 and modified in 1991. Recently we have introduces a two-column procedure that permits quantitation of 65 fatty acids. All three procedures give identical results for three measurements that are the topic of the present report.

It would have been obvious to someone of ordinary skill in the art at the time of the instant invention to modify the lipid composition study of Ruan et al. as applied to claims 2, 3, 5, 6, 12, and 56-58 above in further view of the chromatographic analysis of Moser et al., because while both studies quantify lipids in biological tissues, Moser et al.

has the advantage of using chromatographic analysis of lipids to address peroxisomal disorders.

Rejection #2 under 35 U.S.C. 103(a):

Claims 7, 9, and 13 are rejected under 35 U.S.C. 103(a) as being unpatentable over Ruan et al. as applied to claims 2, 3, 5, 6, 12, and 56-58 above in further view of Watkins et al. [Journal of Lipid Research, volume 39, 1998, pages 1583-1588].

Claim 7 is dependent from claim 5 with the additional limitation of requiring linoleic acid (18:2n6).

Claim 9 is dependent from claim 5 with the additional limitation of requiring cardiolipins.

Claim 13 is dependent from claim 12 with the additional limitation of requiring cardiolipins.

Ruan et al. as applied to claims 2, 3, 5, 6, 12, and 56-58 above does not teach use of cardiolipins and specifically the linoleic acid 18:2 n-6.

The study of Watkins et al., entitled, "Docosahexaxenoic acid accumulates in cardiolipin and enhances HT-29 cell oxidant production," states in the first sentence of the abstract, "The objective of this study was to investigate membrane fatty acids for their effects on mitochondrial function in live cells."

The top of column 2 of page 1583 details some of the specific lipid studied, as stated, "In mammals, CL acyl composition is unusually sensitive to diet, and in humans it is rich in the essential fatty acid linoleic acid (LA, 18:2 n-6)."

It would have been obvious to someone of ordinary skill in the art at the time of the instant invention to modify the lipid composition study of Ruan et al. as applied to claims 2, 3, 5, 6, 12, and 56-58 above in further view of the cardiolipin study of Watkins et al., because while both studies quantify lipids in biological tissues, Watkins et al. has the advantage of quantifying cardiolipins in mitochondria for the purpose of understanding oxidant production and aging.

Rejection #3 under 35 U.S.C. 103(a):

Claims 4 and 14 are rejected under 35 U.S.C. 103(a) as being unpatentable over Ruan et al. as applied to claims 2, 3, 5, 6, 12, and 56-58 above in further view of Watkins et al. in further view of Siguel [US Patent 5,075,101; IDS of 1/5/2004].

Claims 4 and 14 are dependent from claims 3 and 12, respectively, limiting the metabolites to **5,8,11-eicosatrienoic acid, 5,8,11,14,17-eicosapentaenoic acid, 5,8,11-eicosatrienoic acid, and 5,8,11,14,17-eicosapentaenoic acid.**

While Ruan et al. as applied to claims 2, 3, 5, 6, 12, and 56-58 above in view of Watkins et al., teach the method of fatty acid analysis of cells, including EPA, they do not teach the specific nomenclature of the claim required in claim language. Additionally, the above-mentioned prior art does not teach 5,8,11-eicosatrienoic acid (Mead acid).

A web site [www.pdrhealth.com] clarifies the nomenclature and structure of EPA as 5,8,11,14,17-eicosapentaenoic acid.

The patent of Siguel, entitled, "Method of diagnosis of fatty acid or lipid abnormalities," states in column 3, lines 55-65, that Mead acid is an essential fatty acid important in preventing essential fatty acid deficiency.

It would have been obvious for someone of ordinary skill in the art at the time of the instant invention to practice Ruan et al. as applied to claims 2, 3, 5, 6, 12, and 56-58 above in further view of Watkins et al. in further view of Siguel because Siguel shows the advantage of Mead acid in that adequate amounts of Mead acid are required to prevent lipid deficiency in the blood.

Rejection #4 under 35 U.S.C. 103(a):

Claims 17 is rejected under 35 U.S.C. 103(a) as being unpatentable over Ruan et al. as applied to claims 2, 3, 5, 6, 12, and 56-58 above in further view of "The World of Membrane Lipids," [www.biochem.Missouri.edu/~lesa/LIPIDS/membrane_lipid.html]; accessed on 6 December 2006, page made on 2 February 1999].

Claim 17 is dependent from claim 2 wherein displaying generates a web page for viewing.

However, Ruan et al. as applied to claims 2, 3, 5, 6, 12, and 56-58 above does not teach the use of a web page for electronically displaying of results.

"The World of Membrane Lipids," states in its introduction:

This website is an unofficial home for membrane lipid crystal structures. Here, you'll be able to find information about the nomenclature, crystallization, etc. of membrane lipids. Although about 50 structures are known, most of them are not in a database, so the only source of their coordinates is the original journal article.

The purpose of this site is to make this information available to anyone interested, especially structural biologists. To facilitate their use, all coordinate files are in PDB format. If you have any comments or contributions, please send them to Lesa Beamer.

It would be obvious to someone of ordinary skill in the art at the time of the instant invention to modify the lipid quantification method of Ruan et al. as applied to claims 2, 3, 5, 6, 12, and 56-58 above in further view of the web posting database of "The World of Membrane Lipids," because posting lipid results on a web page has the advantage of making the data available to the general public.

Rejection #5 under 35 U.S.C. 103(a):

Claim 10 is rejected under 35 U.S.C. 103(a) as being unpatentable over Ruan et al. as applied to claims 2, 3, 5, 6, 12, and 56-58 above in further view of Grav et al. [Journal of Chromatography B, volume 658, 1994, pages 1-10].

Claim 10 is dependent from claim 6 further limiting the types of internal standards to be employed.

Ruan et al. as applied to claims 2, 3, 5, 6, 12, and 56-58 above does not teach the specific internal standards to be used.

The article of Grav et al., entitled, "Gas chromatographic measurement of 3- and 4-thia fatty acids incorporated into various classes of rat liver lipids during feeding experiments," states in the first sentence of the abstract, "A practical procedure is described for the quantitative measurement of the amount of acyl units derived from tetradecylthioacetic acid (effecting hypolipemia in rats) and tetradecylthiopropionic acid (effecting hyperlipemia)."

The abstract of Grav et al. continues, "The overall recoveries of heptadecanoyl lipids added as internal standards using extraction were 94% to 96%, except for cholesteryl heptadecanoate..."

In Grav et al., section 2.3 on page 2, Grav et al. disclose that one of the species used in claim 10, diheptadecanoyl phosphatidylcholine is used as an internal standard.

It would be obvious to someone of ordinary skill in the art at the time of the instant invention to modify the lipid quantification method of Ruan et al. as applied to claims 2, 3, 5, 6, 12, and 56-58 above in further view of Grav et al. because while Grav et al. disclose a method of quantifying lipids in livers, Grav et al. has the advantage of using the required internal standards in a direct health application in examining hypolipemia and hyperlipemia.

Rejection #6 under 35 U.S.C. 103(a):

Claim 8 is rejected under 35 U.S.C. 103(a) as being unpatentable over Ruan et al. as applied to claims 2, 3, 5, 6, 12, and 56-58 above in further view of Dutta et al. [JAOCS, volume 74, no. 6, 1997, pages 647-657].

Claim 8 is dependent from claim 5 further limiting the types of sterols to be employed as lipids.

Ruan et al. as applied to claims 2, 3, 5, 6, 12, and 56-58 above does not teach use of cholestan-3 β -ols.

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The article of Dutta et al., entitled, "Studies of phytosterol oxides: I: Effect of storage on the content in potato chips prepared in different vegetable oils," states in the abstract:

Potato chips fried in palm oil, sunflower oil, and high-oleic sunflower oil were studied for the content of different phytosterol oxides during 0 to 25 weeks of storage in the dark. Oxidation products of sitosterol (2,4 alpha-ethyl-5-cholesten-2b-ol) and campesterol (2,4 alpha methyl-5cholesten-3b-ol) were synthesized to help identify the phytosterol oxides.

Dutta et al. continue in the introduction to explain in the first sentence of the introduction:

Abundant information exists on the formation of cholesterol oxidation products in foods and their biological implications, but there is relatively little on such products generated from phytosterols.

It would have been obvious to someone of ordinary skill in the art at the time of the instant invention to modify the lipid quantification method of Ruan et al. as applied to claims 2, 3, 5, 6, 12, and 56-58 above in view of the phytosterol quantitation method of Dutta et al. because the study of Dutta et al. has the advantage of using the required fatty acids for further understanding of biological implications of cholesterol and phytosterols.

Conclusion

Papers related to this application may be submitted to Technical Center 1600 by facsimile transmission. Papers should be faxed to Technical Center 1600 via the central PTO Fax Center. The faxing of such pages must conform with the notices published in the Official Gazette, 1096 OG 30 (November 15, 1988), 1156 OG 61 (November 16, 1993), and 1157 OG 94 (December 28, 1993)(See 37 CFR § 1.6(d)). The Central PTO Fax Center Number is (571) 273-8300.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Russell Negin, Ph.D., whose telephone number is (571) 272-1083. The examiner can normally be reached on Monday-Friday from 7am to 4pm.

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If attempts to reach the examiner by telephone are unsuccessful, the examiner's Supervisor, Ram Shukla, Supervisory Patent Examiner, can be reached at (571) 272-0735.

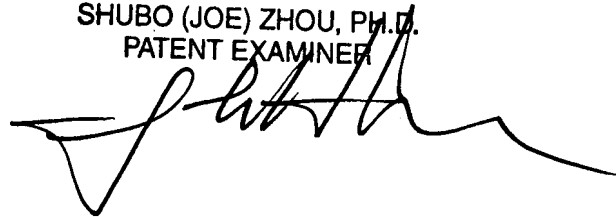
Information regarding the status of the application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information on the PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

RSN

8 June 2007

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SHUBO (JOE) ZHOU, PH.D.
PATENT EXAMINER

A handwritten signature in black ink, appearing to be 'Shubo Zhou', written over the printed name and title.